Under the Paperwork Reduction Act of 1995, no persons are required to respo

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number		10595453		
Filing Date		2006-04-20		
First Named Inventor Jame		s W. Larrick		
Art Unit			-	
Examiner Name				
Attorney Docket Number	er.	53233-00008		

					U.S.	PATENTS			Remove	
Examiner Initial*	Cite No	Patent Number	Kind Code ¹	Issue E	Date	Name of Patentee or Applicant of cited Document		Pages,Columns,Lines when Relevant Passages or Rele- Figures Appear		
	1	6569441	B1	2003-0	5-27	Kunz et al.				
If you wisl	h to a	dd additional U.S. Pate	nt citatio	n inform	ation pl	ease click the	Add button.		Add	
			U.S.P	ATENT	APPLI	CATION PUBL	LICATIONS		Remove	
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	tion	Name of Patentee or Applicant of cited Document		Releva	Columns,Lines where nt Passages or Relev Appear	
	1	20020165268	A1	2002-1	1-07	Wechter				
If you wis	h to a	dd additional U.S. Publ	shed Ap	plication	citatio	n information p	lease click the Ad	d button.	Add	
				FOREIG	SN PAT	ENT DOCUM	ENTS		Remove	
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²		Kind Code ⁴	Publication Date	Name of Patente Applicant of cited Document	e or w	'ages,Columns,Lines /here Relevant 'assages or Relevant 'igures Appear	+=
	1									
If you wis	h to a	dd additional Foreign P	atent Do	cument	citation	information pl	ease click the Add	button	Add	_
			NON	-PATE	NT LITE	RATURE DO	CUMENTS		Remove	
Examiner Cite Initiats* Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, pages(s), volume-issue number(s), publisher, city and/or country where published.									Тs	

	Application Number		10595453	
NEODMATION DIOOLOGUEE	Filing Date		2006-04-20	
NFORMATION DISCLOSURE STATEMENT BY APPLICANT	First Named Inventor Jame		es W. Larrick	
Not for submission under 37 CFR 1.99)	Art Unit			
Not for Submission under or or it 1.557	Examiner Name			
	Attorney Docket Numb	er	53233-00008	

	1			
If you wish	h to a	d additional non-patent literature document citation information please click the Add button	Add	

EXAMINER SIGNATURE

EXAMINER SIGNATURE

Examiner Signature

Date Considered

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

See Kint Code of USPTO Patent Documents at invenUSPTO_GDI/ or MPEP 901.04. * Enter of thice that issued the document, by the involved ow (WIPO Standard ST3.). * For Suppresse patent counters, the noticeation of the year of the region of the Emperor many precede the sent annual replication of the part document.

Ind of counter by the appropriate symbols as endicated on the document under WIPO Standard ST1.6 if possole, **Applicant is to place a check mark here if Emploit languages translation is attached.

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99) Art Unit

Application Number		10595453			
Filing Date		2006-04-20			
First Named Inventor Jame		s W. Larrick			
Art Unit					
Examiner Name					
Attornou Docket Numb	nr.	52222 00000			

CERTIFICATION STATEMENT

Please see	37	CFR :	1 97	and	1 98	to make	the	appropriate	selection(s)	

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information siclosure statement. See 37 CFR 197(e)(1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 15(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(eV).

- 7 See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- □ None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/C. Rachal Winger/	Date (YYYY-MM-DD)	2006-07-17
Name/Print	C. Rachal Winger	Registration Number	55815

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 GA 37 CFR.

1.14. This collection is estimated to take I hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, u.S. Operatment of Commence, P. O. Box 1430, Alexandriu, V.S. 2213.1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandriu, V.S. 2213.1450.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is \$3 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2504 and 2506. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the state of the s
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.